Listing of Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1. (Previously Presented) An isolated tribonectin comprising a boundary-lubricating amount of a polypeptide, said polypeptide comprising the amino acid sequence of SEQ ID NO:1 and at least one O-linked oligosaccharide moiety, wherein the molecular weight of said tribonectin is in the range of 220-280 kDa.
- 2. (Previously Presented) The tribonectin of claim 1, wherein said moiety is a $\beta(1-3)$ Gal-GalNAc moiety.

3. - 9. (Cancelled)

- 10. (Previously Presented) The tribonectin of claim 1, wherein said O-linked oligosaccharide moiety of said polypeptide reduces the coefficient of friction between bearing surfaces.
- 11. (Currently Amended) The tribonectin of claim 1, wherein said O-linked oligosaccharide moiety of said tribonectin reduces the coefficient of friction between bearing surfaces *in vitro*.
- 12. (Currently Amended) The tribonectin of claim 1, wherein said O-linked oligosaccharide moiety of said tribonectin reduces the coefficient of friction between bearing surfaces *in vivo*.
- 13. (Previously Presented) The tribonectin of claim 1, wherein addition of said tribonectin to a solution does not increase the viscosity of said solution by more than 10%.

14-15. (Cancelled)

- 16. (Previously Presented) The tribonectin of claim 1, wherein at least 10% of said tribonectin is glycosylated by said O-linked oligosaccharide moiety.
- 17. (Previously Presented) The tribonectin of claim 1, wherein at least 40% of said tribonectin is glycosylated by said O-linked oligosaccharide moiety.
 - 18. 39. (Cancelled)
- 40. (Previously Presented) A biocompatible composition comprising the isolated tribonectin of claim 1, wherein said composition is in the form of a film, membrane, foam, gel, or fiber.
 - 41. 54. (Cancelled)
- 55. (Previously Presented) The tribonectin of claim 1, further comprising hyaluronic acid.
- 56. (Currently Amended) A composition comprising a boundary-lubricating polypeptide encoded by a nucleic acid <u>construct</u>, <u>said construct comprising a human megakaryocyte stimulating factor coding sequence</u>, wherein <u>said megakaryocyte stimulating factor coding sequence consists eonsisting essentially</u> of exon 1, 2, 3, 4, and 6-12 of a human megakaryocyte stimulating factor gene <u>and lacks at least one exon of said megakaryocyte stimulating factor gene</u>.
 - 57. (Cancelled)
- 58. (Currently Amended) A composition comprising a boundary-lubricating polypeptide encoded by a nucleic acid construct, said construct comprising a human megakaryocyte stimulating factor coding sequence, wherein said megakaryocyte stimulating factor coding sequence consists consisting essentially of exon 1, 3, and 6-12 of a human

megakaryocyte stimulating factor gene and lacks at least one exon of said megakaryocyte stimulating factor gene.

59. (Currently Amended) A composition comprising a boundary-lubricating polypeptide encoded by a nucleic acid <u>construct</u>, <u>said construct comprising a human megakaryocyte stimulating factor coding sequence</u>, wherein said megakaryocyte stimulating <u>factor coding sequence consists consisting essentially</u> of exon 1 and 6-12 of a human megakaryocyte stimulating factor gene <u>and lacks at least one exon of said megakaryocyte</u> stimulating factor gene.

For example, claim 56 requires a polypeptide encoded by a construct containing a megakaryocyte stimulating factor coding sequence consisting of exons 1, 2, 3, 4, and 6-12 and lacking at least one exon of the human megakaryocyte stimulating factor gene. Claim 58 requires a polypeptide encoded by a construct containing a megakaryocyte stimulating factor coding sequence consisting of exons 1, 3, and 6-12 and lacking at least one exon of the human megakaryocyte stimulating factor gene. Finally, claim 59 requires a polypeptide encoded by a construct containing a megakaryocyte stimulating factor coding sequence consisting of exons 1, and 6-12 and lacking at least one exon of the human megakaryocyte stimulating factor gene. In view of this amendment, Applicant submits that the claims no longer read on Turner's description of a full-length (12 exons) megakaryocyte stimulating factor gene.

Withdrawal of this rejection is respectfully requested.